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# Genetic risk factors for type 2 diabetes mellitus and response to sulfonylurea treatment

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**Objective** After the identification of type 2 diabetes mellitus (T2DM) risk alleles from genome-wide association studies, models have been developed to identify subjects at high risk to develop T2DM. We hypothesize that a panel of 20 repeatedly associated T2DM risk alleles influences response to sulfonylureas (SUs).

**Methods** Two hundred and seven incident SU (tolbutamide, glibenclamide, glimepiride, gliclazide) users with T2DM were recruited from four primary care centers. A genetic risk score per patient was calculated based on the number of risk-alleles. With this score, patients were categorized into three predefined genetic risk groups. The effect of the genetic risk group on the achievement of stable SU dose, prescribed stable SU dose, and time to stable SU dose was analyzed.

**Results** Carriers of more than 17 T2DM risk alleles had a 1.7-fold reduced likelihood to achieve stable SU dose ( $P=0.044$ ). No significant effect of the number of T2DM risk alleles on prescribed dose was found. Carriers of more than 17 T2DM risk alleles showed a marginally significant

increased time to stable dose (hazard ratio: 0.81; 95% confidence interval, 0.75–1.01,  $P=0.058$ ).

**Conclusion** T2DM risk alleles are associated with response to SUs in primary care T2DM patients. This suggests that individualization of T2DM treatment according to genetic profile may be an opportunity to improve clinical outcome. *Pharmacogenetics and Genomics* 21:461–468 © 2011 Wolters Kluwer Health | Lippincott Williams & Wilkins.

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**Keywords:** glibenclamide, gliclazide, glimepiride, pharmacogenetics, sulfonylureas, tolbutamide, type 2 diabetes mellitus

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## Introduction

The incidence of type 2 diabetes mellitus (T2DM) is increasing at an alarming rate. Worldwide, the number of patients is expected to increase from 171 million in 2000 to 366 million in 2030 [1]. The therapeutic goal of treating T2DM patients is to prevent or delay long-term microvascular and macrovascular complications by achieving the best possible glycemic control.

Sulfonylureas (SUs) are part of the mainstay of treatment with oral antidiabetic drugs. Tolbutamide, glibenclamide (glyburide), gliclazide, and glimepiride are the most commonly used representatives of this group. These drugs act by closing the pancreatic  $\beta$ -cell potassium channels, stimulating insulin secretion [2]. SUs are initiated at a low dose and escalated to the optimal dose with intervals of 2–4 weeks until the glycemic target ( $\text{HbA1c} < 7\%$ ) is achieved. However, there is significant interpatient variability in response to SUs, with approximately 10–20% of the patients experiencing primary failure (decrease in fasting glucose level  $< 1.1 \text{ mmol/l}$ )

and a similar percentage having an above average response (mean reduction  $\text{HbA1c}$  1.5–2%) [3–5].

With the completion of multiple genome-wide association studies (GWAS) the knowledge of the complex genetic background of T2DM has increased. These studies report associations between genetic variants and the risk for the development of T2DM. A panel of 20 T2DM associated single nucleotide polymorphisms (SNPs) comprising 19 genes out of the GWAS data appears, that has been replicated in several studies [6–17]. These SNPs are used in models with the ultimate goal to identify subjects at high risk to develop T2DM. Although marginally, the addition of genetic information to clinical T2DM risk factors increased the ability to predict future diabetes [18–24].

From the panel of 20 T2DM risk-associated SNPs, the majority is involved in the process of insulin release from the pancreatic  $\beta$ -cells (Table 1). As SUs act by stimulating insulin secretion, response to SU treatment may also be influenced by these genetic variants. Indeed, two of the 19 T2DM risk-associated genes, encoding *KCNJ11* and *TCF7L2*, have been previously correlated with variation in SU response [4]. Furthermore, in subjects

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**Table 1 Selected single nucleotide polymorphisms associated with type 2 diabetes mellitus**

Gene	rs number	Chromosome	Risk allele	Year	Mechanism	References
<i>NOTCH2</i>	rs10923931	1	T	2008	Unknown	[9,19–24]
<i>THADA</i>	rs7578597	2	T	2008	Unknown	[9,19–24]
<i>IGF2BP2</i>	rs4402960	3	T	2007	β-cell dysfunction	[8,9,11–13,18–24]
<i>PPARG</i>	rs1801282	3	C	2000	Insulin sensitivity	[8,9,11–13,19–24]
<i>ADAMTS9</i>	rs4607103	3	C	2008	Unknown	[9,19–24]
<i>WFS1</i>	rs10010131	4	G	2007	Unknown	[9,13,18–20,22,24]
<i>CDKAL1</i>	rs7754840	6	C	2007	β-cell dysfunction	[7–9,11–13,18–24]
<i>JAZF1</i>	rs864745	7	A	2008	β-cell dysfunction	[9,19–24]
<i>SLC30A8</i>	rs13266634	8	C	2007	β-cell dysfunction	[7–9,11–14,18–24]
<i>CDKN2A/CDKN2B</i>	rs10811661	9	T	2007	β-cell dysfunction	[8,9,11–13,18–24]
	rs564398	9	A			[8,18,20,24]
<i>TCF7L2</i>	rs7903146	10	T	2006	β-cell dysfunction	[7–9,11–14,18–24]
<i>HHEX/IDE</i>	rs1111875	10	G	2007	β-cell dysfunction	[7–9,11–14,18–24]
<i>CDC123/CAMK1D</i>	rs12779790	10	G	2008	Unknown	[9,19–24]
<i>KCNJ11</i>	rs5219	11	T	2003	β-cell dysfunction	[8,9,11–13,19–24]
<i>KCNQ1</i>	rs2237892	11	C	2008	β-cell dysfunction	[10,17,24]
<i>MTNR1B</i>	rs10830963	11	G	2009	Disturbance of circadian rhythm	[6,16,24]
<i>TSPAN8/LGR5</i>	rs7961581	12	C	2008	Unknown	[9,19–24]
<i>FTO</i>	rs8050136	16	A	2007	Obesity	[8,9,11,13,19–22,24]
<i>HNF-1β (TCF2)</i>	rs757210	17	A	2007	β-cell dysfunction	[15,20,21,24]

analyzed for genetic variation in the genes *TCF7L2*, *PPARG*, *FTO*, *KCNJ11*, *NOTCH2*, *WFS1*, *CDKAL1*, *IGF2BP2*, *SLC30A8*, *JAZF1*, *HHEX*, it was reported that subjects with 12 or more T2DM risk alleles did not increase their insulin secretion to compensate for the increased insulin resistance as efficiently as those with 8 or less risk alleles [19]. Therefore, patients with a greater number of risk alleles may show less response to SU treatment and individualization of T2DM treatment according to genetic profile may be an opportunity to improve clinical outcome.

We hypothesize that the genetic variants associated with the development of T2DM are also associated with response to SU treatment. Therefore, we investigated the effect of T2DM risk alleles on the response to SU treatment in T2DM patients in a primary care setting.

## Methods

### Study setting

In the Netherlands the general practitioner (GP) plays a central role in the provision of health care. Patients are listed with one GP who is consulted for all healthcare problems and indicates whether a referral to secondary care is appropriate. Typically, the GP keeps an electronic patient record (EPR) that covers all medical information concerning the patient including prescription information and reports from laboratories and specialists. GP's have adopted the practice guideline T2DM of the Dutch College of General Practitioners [25]. Tailoring the treatment to the individual patient is an important part of the therapy.

### Cohort ascertainment

A total of 207 T2DM patients from four university-affiliated primary care centers (17 GPs) located in the vicinity of Leiden, the Netherlands were recruited. The ascertainment of the cohort has been described in detail

previously [26]. In brief, patients that had at least one prescription of tolbutamide, glibenclamide, glimepiride, or gliclazide between January 1992 and June 2008, were at least 18 years of age and without insulin use at the time of first SU prescription, and had at least 270 days of follow-up registered in the EPR, were included. Ethnicity was not routinely recorded in the EPR but most patients in the Netherlands are from European ancestry. Patients received a written invitation by mail from their GP. Of the 472 invited patients, 222 (47%) agreed to participate (see Fig. S1, Supplemental digital content 1, <http://links.lww.com/FPC/A263>, cohort ascertainment). After consent, a saliva collection kit (DNA Genotek Inc., Ottawa, Ontario, Canada) was mailed. The study was approved by the ethics committee of the Leiden University Medical Center.

### Genotyping

We selected a panel of 20 SNPs in 19 genes that have been associated with the development of T2DM in at least three GWAS and were consistently replicated in later studies aimed at estimating the predictive value of these SNPs on the development of T2DM [6–24]. The selected SNPs are listed in Table 1. DNA was isolated from the saliva according to the protocol provided by the manufacturer (DNA Genotek Inc.). Taqman genotyping assays for 19 SNPs were designed by and obtained from Applied Biosystems (Applied Biosystems, Nieuwerkerk aan den IJssel, the Netherlands). SNP rs757210 could not be designed as a Taqman genotyping assay and therefore was genotyped by pyrosequencing (Isogen Life Science, Maarssen, the Netherlands). Taqman genotyping assays were performed on the LightCycler 480 II Real-Time PCR System (Roche Diagnostics, Almere, the Netherlands) according to standard procedures. Genotyping was performed without knowledge of the clinical data. We obtained an average genotyping success rate of more than 95%. As a quality control 5% of the samples were

genotyped in duplicate for all assays and no inconsistencies were observed. Five patients were excluded for quality reasons (genotype call rate  $\leq 80\%$ ). All SNPs were in Hardy–Weinberg equilibrium ( $P > 0.05$ ), with the exception of rs2237892 ( $P = 0.011$ ). This is most probably ascribed to the very low minor allele frequency of rs2237892, which was 0.025 in our study and comparable with previously reported minor allele frequencies of 0.056–0.075 ([http://www.ncbi.nlm.nih.gov/projects/SNP/snp\\_ref.cgi?rs=2237892](http://www.ncbi.nlm.nih.gov/projects/SNP/snp_ref.cgi?rs=2237892)) accessed 5 October 2010.

### Definition of effect

For each patient a cumulative genetic risk score was calculated based on the number of present risk alleles. Each person could have 0, 1, or 2 of them for each SNP, resulting in a theoretical individual cumulative risk score between 0 and 40. This approach assumes an equal and additive effect of each allele on the risk of T2DM. To allow categorization of patients, we predefined three genetic risk groups on the basis of the frequency distribution of risk alleles. We defined a low genetic risk group and a high genetic risk group as the quintiles with the lowest and highest number of T2DM risk alleles, respectively. All other patients (three quintiles) were categorized in the intermediate risk group.

The primary endpoint of our study is the effect of the genetic risk group on achieving stable SU dose. Stable SU dose was defined as the first period of more than or equal to 270 consecutive days without SU dose adjustment, or initiation or adjustment of therapy with other SUs, insulin or metformin. If therapy with insulin was initiated patients were censored. The period of more than or equal to 270 days was chosen because prescriptions in the Netherlands are limited to a maximum of 90 days and more than or equal to 270 days equals three consecutive prescriptions. Stable SU dose was calculated as normalized dose by dividing the prescribed daily dose with the standard daily dose used by the Pharmaceutical Aid Committee of the Dutch Health Care Insurance Board (10 mg glibenclamide; 1000 mg tolbutamide; 160 mg gliclazide; 2 mg glimepiride). Secondary endpoints of our study are the stable SU dose and the time required for dose escalation (time to stable SU dose).

### Statistical analysis

The data were analyzed using the SPSS statistical package (version 16.0, SPSS, Chicago, Illinois, USA). Deviation from Hardy–Weinberg equilibrium was tested by the  $\chi^2$  test. Achievement of stable SU dose was analyzed with the  $\chi^2$  test and multivariate logistic regression analysis. Differences in mean stable SU dose between genetic risk groups were analyzed using the Kruskal–Wallis test and multivariate linear regression analysis. Associations between the genetic risk groups and time to stable SU dose were evaluated using the Cox survival regression analysis. Before multivariate analysis,

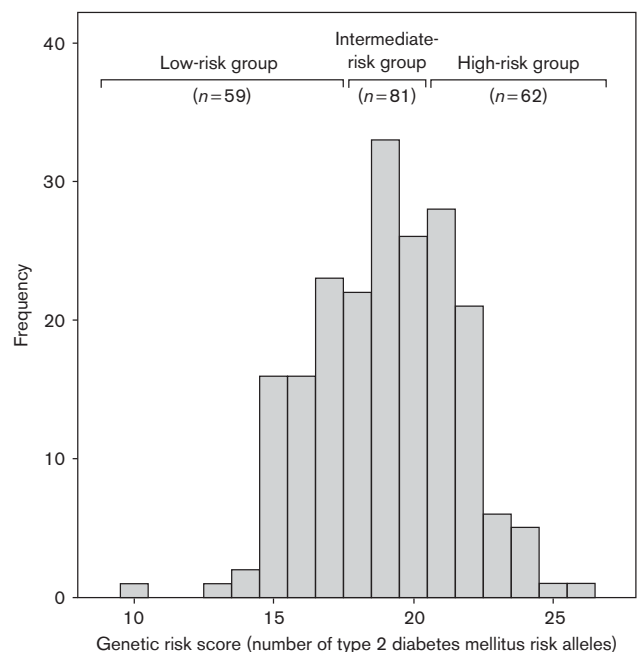
all demographic and clinical variables were tested univariately against the selected outcome. Variables with a  $P$  value of 0.1 or less, age, and sex were selected for multivariate analysis. All multivariate analyses were corrected for age and sex.

## Results

Data from 202 T2DM patients were available. The range of the calculated genetic risk score was 10–26. The quintiles with the lowest ( $\leq 17$ ) and highest ( $\geq 21$ ) number of T2DM risk alleles consisted of 59 patients and 62 patients, respectively (Fig. 1). Table 2 presents the characteristics of the 202 patients. There were no differences between the different genetic risk groups observed in any of the patient characteristics except for age. Patients in the high-risk group were younger at the time of first SU prescription compared with patients in the low-risk and intermediate-risk group, respectively ( $P = 0.001$ ). Mean follow-up was 5.9 years, reflecting that most patients (75.2%) were included after 2000. Our patients received an average of 26 SU prescriptions during the follow-up period with a median duration of 90 days per prescription.

The results of achieving stable SU dose and the T2DM genetic risk groups are presented in Fig. 2. Of the patients, 148 (73.3%) achieved stable SU dose. The percentage of

Fig. 1



Distribution of type 2 diabetes mellitus risk alleles and subsequent classification in risk groups. Patients were categorized in three genetic risk groups. Low-risk group; quintile with the lowest ( $\leq 17$ ) number of T2DM risk alleles. High-risk group; quintile with the highest ( $\geq 21$ ) number of T2DM risk alleles. Remaining patients were categorized in the intermediate-risk group (18–20 T2DM risk alleles).

**Table 2** Characteristics of the 202 patients with type 2 diabetes mellitus in primary care

Variable no. (%) <sup>a</sup>	All patients	Genetic risk group			<i>P</i> value
		Low-risk	Intermediate-risk	High-risk	
Subjects	202	59 (29.2)	81 (40.1)	62 (30.7)	NA
Men	106 (52.5)	30 (50.8)	45 (55.6)	31 (50.0)	0.77
Women	96 (47.5)	29 (49.2)	36 (44.4)	31 (50.0)	
Age in years, mean (SD)	61.4 (10.7)	64.0 (9.5)	62.6 (10.4)	57.3 (11.1)	<b>0.001</b>
Follow-up in years, mean (SD)	5.9 (3.0)	6.0 (3.0)	5.7 (3.0)	6.2 (3.0)	0.52
Visits in year one (SD)	9.6 (4.7)	8.6 (3.7)	10.0 (5.2)	10.0 (4.6)	0.35
Metformin use	62 (30.7)	18 (30.5)	27 (33.3)	17 (27.4)	0.75
Primary sulfonylurea					0.098 <sup>b</sup>
Glibenclamide	12 (5.9)	7 (11.9)	1 (1.2)	4 (6.5)	
Tolbutamide	85 (42.1)	18 (30.5)	41 (50.6)	26 (41.9)	
Gliclazide	24 (11.9)	7 (11.9)	10 (12.3)	7 (11.3)	
Glimepiride	81 (40.1)	27 (45.8)	29 (35.8)	25 (40.3)	

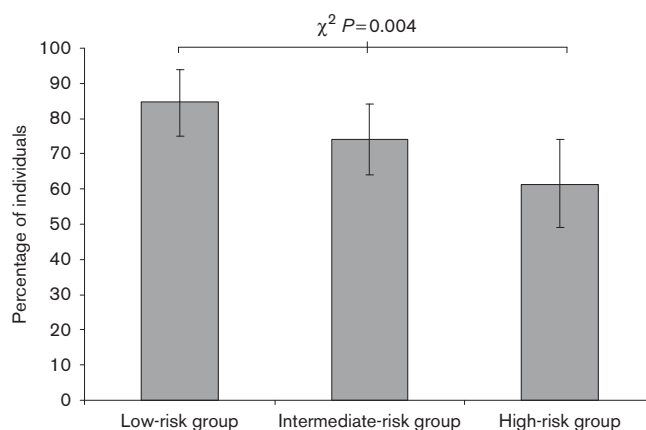
Low-risk group; patients with 17 or less risk alleles. Intermediate-risk group: patients with 18–20 risk alleles. High-risk group; patients with at least 21 risk alleles.

NA, not applicable; SD, standard deviation.

<sup>a</sup>Unless stated otherwise.

<sup>b</sup> $\chi^2$  for primary sulfonylurea vs. genetic risk group.

*P* value < 0.05 is regarded as significant and indicated with bold font.

**Fig. 2**

Percentage of type 2 diabetes mellitus patients that reached stable sulfonylurea dose. Low-risk group; patients with 17 or less risk alleles. Intermediate-risk group: patients with 18–20 risk alleles. High-risk group; patients with at least 21 risk alleles.

patients achieving stable SU dose was lower in the high-risk group compared with the intermediate-risk and low-risk groups (61.3 vs. 74.1 vs. 84.7%, respectively,  $P = 0.004$ ). In the multivariate logistic regression analysis age at first SU prescription, the concomitant use of metformin, and the T2DM genetic risk group were independently significantly associated with achieving stable SU dose (Table 3). The regression model explained 28.7% of the variation in achievement of stable dose. Data show that patients with a higher T2DM risk had a 1.7-fold reduced likelihood to achieve stable SU dose ( $P = 0.044$ ).

Next, the mean SU starting dose was analyzed. The mean SU starting dose for all patients was 0.61 (95% CI: 0.58–0.65). As expected, no differences in SU starting dose were found between the different genetic risk groups. No differences in mean stable SU dose were found between

the different T2DM genetic risk groups [low-risk group 0.90, 95% confidence interval (CI) 0.75–1.05 vs. intermediate-risk group 0.84, 95% CI: 0.74–0.94 vs. high-risk group 0.95, 95% CI: 0.72–1.17,  $P = 0.97$ ]. In multivariate linear regression, only the effect of the SU starting dose and sex were independently significant associated with stable SU dose, whereas the genetic risk group for T2DM was not associated with stable SU dose.

As SUs are escalated to the optimal dose, the effect of the genetic risk group on time to stable SU dose was evaluated. Carriers of the high-risk genetic profile ( $\geq 21$  risk alleles) had a two-fold and five-fold longer time to stable dose compared with patients with the intermediate risk (18–20 risk alleles) and low-risk profile ( $\leq 17$  risk alleles) (median time to stable SU dose 160 vs. 59 vs. 31 days, respectively,  $P = 0.007$ ). In a multivariate Cox regression analysis including the factors such as age

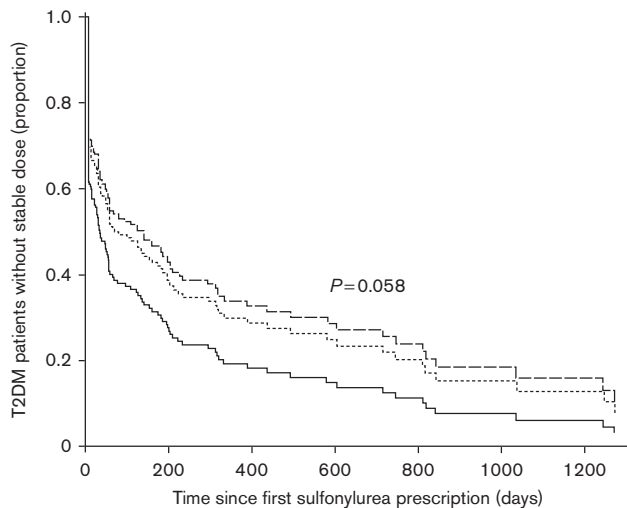
**Table 3** Analysis of factors relevant for achieving stable sulfonylurea dose in patients with type 2 diabetes mellitus in primary care

Factor	Univariate				Multivariate <sup>a</sup>		
	OR	95% CI	R <sup>2</sup>	P value	OR	95% CI	P value
Constant					1.81	NA	0.654
Male vs. female sex	1.55	0.83–2.91	0.009	0.17	1.54	0.72–3.29	0.262
Effect of age (per year increase)	1.06	1.03–1.09	0.069	<b>&lt;0.001</b>	1.04	1.00–1.08	<b>0.036</b>
Metformin use vs. no metformin use at stable sulfonylurea dose	0.07	0.03–0.16	0.238	<b>&lt;0.001</b>	0.07	0.03–0.17	<b>&lt;0.001</b>
Genetic risk group (low-risk → intermediate-risk → high-risk group)	0.54	0.35–0.82	0.042	<b>0.004</b>	0.59	0.35–0.99	<b>0.044</b>

→, stepwise increase from low-risk to intermediate-risk to high-risk group; CI, confidence interval; OR, odds ratio; R<sup>2</sup>, variation in the achievement of stable dose.

<sup>a</sup>R<sup>2</sup> for the complete model was 0.287.

P value <0.05 is regarded as significant and indicated with bold font.

**Fig. 3**

Multivariate Cox regression analysis plots of time to the first stable dose of sulfonylureas in type 2 diabetes mellitus patients in primary care. Low-risk group (—); patients with 17 or less risk alleles. Intermediate-risk group (---); patients with 18–20 risk alleles. High-risk group (···); patients with at least 21 risk alleles.

on first SU prescription, sex, and the concomitant use of metformin, patients with a higher number of risk alleles showed a marginally significant increased time to stable SU dose (hazard ratio: 0.81; 95% CI: 0.75–1.01,  $P = 0.058$ ) (Fig. 3).

## Discussion

To the best of our knowledge, this is the first study exploring the relationship between response to treatment with SUs and T2DM risk alleles. In this retrospective cohort study of 202 T2DM patients, patients with more than 17 risk alleles have a 1.7-fold reduced likelihood to achieve a stable SU dose. These patients also show a marginally significant increased time to achieve stable SU dose compared with carriers of less than 17 risk alleles. However, the number of T2DM risk alleles does not seem to affect the average stable SU dose used. Therefore, our data suggest that patients with a higher number of T2DM risk alleles have a decreased and delayed response to SU treatment.

Drug response is determined by both pharmacokinetics and pharmacodynamics of a drug. Several groups have investigated genetic variation in genes affecting the pharmacokinetics of SU response. Two variants in *CYP2C9*, *CYP2C9\*2* and *CYP2C9\*3*, have been associated with a decreased SU metabolism in healthy volunteers [27]. Five studies assessed the effect of these polymorphisms in T2DM patients. Presence of the *CYP2C9\*3* allele was associated with an increased risk for hypoglycemia [28,29]. Tolbutamide users with a *CYP2C9\*2* or *CYP2C9\*3* allele have been shown to have a significantly lower dose escalation compared with homozygous carriers of the *CYP2C9\*1* allele [30]. In a large cohort of 1073 incident SU users with T2DM Zhou *et al.* [31] found that carriers of the *CYP2C9\*2* or *CYP2C9\*3* allele were less likely to fail on SU monotherapy. In a recent study we found no statistically significant effect of the *CYP2C9\*2* or *CYP2C9\*3* allele on the prescribed stable dose [26].

Variation in genes associated with the pharmacodynamics of SUs in T2DM patients has received considerably less attention. Genetic variants associated with SU response have been described for some monogenic forms of diabetes [32–34]. For polygenic T2DM, variants in the genes *KCNJ11*, *TCF7L2*, *ABCC8*, *IRS1*, and *NOS1AP* have been associated with SU response [35–38]. Of these, only the genes *KCNJ11* and *TCF7L2* were reported to contribute to an increased risk for T2DM in published GWAS. *KCNJ11* encodes the Kir6.2 subunit, one of the two subunits that form the ATP-sensitive potassium channel involved in insulin release. Carriership of the E23K variant of the *KCNJ11* gene has been associated with failure to SU therapy, but there are some conflicting results [39–41]. Variants in the *TCF7L2* gene have also been associated with SU response. In a study with 901 incident SU users, patients with the TT genotype for rs7903146 were 1.73 times less likely to be treated to lower a target HbA1c of 7% in the first 3–12 months of treatment compared with patients with the CC genotype [42]. For a variant in linkage with rs7903146 an even larger effect (odds ratio = 1.95) was reported. In this study, none of the individual risk alleles were significantly associated with the achievement of stable dose (see Table S2, Supplemental digital content 2, <http://links.lww.com/FPC/A264>), risk allele frequency and association with

achievement of stable SU dose of the individual SNPs). This is most likely due to the limited sample size of our study and the probable small effect size of the individual risk alleles.

Our study has some limitations. No data were available for patients that switched to another GP or who died after 1992. Therefore, we cannot completely rule out the possibility of selection bias, although this is conceptually very unlikely. A nonresponse analysis with age, sex, type of first prescribed SU, metformin use, and GP showed no differences between participants and patients who did not consent to our study, suggesting that no selection bias has occurred on any of these parameters.

We selected stable SU dose as the primary endpoint of our analysis. Ideally macrovascular (e.g. diabetes-related death or myocardial infarction) or microvascular events (e.g. retinopathy or renal failure) would have been used. Alternatively, biomarkers related to these events, such as HbA1c or fasting plasma glucose (FPG) might have been used. However, as data concerning these parameters were not routinely recorded in the EPR, data were too sparse to be used in our analysis. Therefore, we selected stable SU dose as an alternative. Although, no SU pharmacogenetics studies have used stable SU dose as endpoint, this parameter closely reflects actual clinical practice. The time to stable SU dose analysis assumes that GPs adhere to the T2DM guideline of the Dutch College of General Practitioners and titrate SU dose in response to glucose and HbA1c levels. We have three arguments that support our assumption. Firstly, mean FPG was 7.77 mmol/l (95% CI: 7.42–8.12,  $n = 95$ ) for the subgroup of patients with a FPG measurement available during stable SU dose. Secondly, the adherence of GPs to guidelines is reported to be good in the Netherlands [43]. Finally, even if GPs do not adhere to the T2DM guideline, and bias would be introduced to our analysis, there is no reason to assume that the nonadherence of GPs is not divided randomly over the different genetic risk groups. Therefore, possible nonadherence does not affect the comparison of the time to stable dose between the different genetic risk groups but can only affect the absolute results of this analysis.

There are multiple known factors that predict a good response to SUs including baseline HbA1c, recently diagnosed diabetes, mild-to-moderate fasting hyperglycemia ( $< 12.2$ – $13.3$  mmol/l), good  $\beta$ -cell function (high fasting C-peptide level), no history of insulin therapy, and absence of islet cell or glutamic acid decarboxylase antibodies [3]. However, for none of these factors sufficient data were available in our retrospective cohort study and we were unable to account for their effect. In addition, the available data on weight, a factor that is associated with the onset of T2DM, were too sparse to be included in the analysis as a covariate. As a consequence we cannot rule out that patients with a higher number of

risk alleles also have a more severe form of T2DM that might confer to an a priori decreased probability to achieve stable SU dose. In our opinion, the only way to collect sufficient high quality data that cover all of these parameters would be to conduct a prospective observational study. Ideally such a study would include two treatment arms with pharmacological different drugs or placebo. Such a design would allow differentiating between the effect of T2DM risk alleles on disease progression and effect on treatment.

The results of different SUs were pooled in one analysis. Although SUs are generally reported to have equipotent glucose lowering effects when administered in maximally effective doses [3,5], it would be interesting to investigate if our hypothesis is valid for each of the individual SUs. However, due to the sample size of our study such a subgroup analysis was not possible.

We achieved a high success rate of genotyping with a call rate of more than 95% for all individual SNPs. After exclusion of five patients with a call rate of less than or equal to 80, 0.9% of the genotype data were missing. Missing genotype data were replaced with a risk score of 0. To test the sensitivity of our analysis for this replacement, we reanalyzed the data using two alternative approaches. First, as for some SNPs the wild-type allele is the risk allele, missing data were replaced with the score of the wild-type allele. As a result, two patients were reclassified from the low-risk to the intermediate-risk group, and one patient was reclassified from the intermediate-risk group to the high-risk group. Secondly, we excluded all patients with any missing data, resulting in the exclusion of an additional 31 (15.3%) patients. Similar results on all end points were obtained with all approaches, except for the effect of the genetic risk score that lost statistical significance in multivariate analysis after exclusion of all patients with missing data. These sensitivity analyses indicate that our results are valid.

The analysis of the effect of the genetic risk score on SU response assumes that each risk allele has an equal and additive effect, both within and between loci. This is clearly a simplification of the mechanism leading to variation of SU response. However, this approach is used in all GWAS studies concerning prediction of T2DM. Until it is clear what the true effect size of individual risk alleles is, the additive genetic model is probably the most appropriate and consistent method to analyze T2DM genetic data.

We chose to compare the quintile with the lowest ( $\leq 17$ ,  $n = 59$ , low-risk group) and highest ( $\geq 21$ ,  $n = 62$ , high-risk group) number of T2DM risk alleles, whereas patients with 18–20 risk alleles were pooled in one group ( $n = 81$ , intermediate-risk group) (Fig. 1). The use of quintiles was based on a study by Lyssenko *et al.* [19] and allows potentially easy translation to the clinic by clear



classification of T2DM patients. The cutoffs for the quintiles with the highest and lowest number of T2DM risk alleles fell within the group of patients with 21 and 17 risk alleles respectively. We categorized patients with 17 risk alleles to the low-risk group and 21 alleles to the high-risk group, resulting in a slightly larger number of patients in both categories than anticipated. To ascertain that our results are not solely due to study design, we also analyzed the genetic risk score as a continuous variable instead of the analysis of risk groups. Next to this genetic risk score (range 10–26), sex, age on first SU prescription, and the use of metformin were included in the multivariate analysis. Data showed similar results for both the effect size and direction for the genetic risk score (odds ratio 0.88 95% CI: 0.76–1.02,  $P = 0.11$ ). This suggests that with increasing number of risk alleles, the chance of achieving stable SU dose decreases.

The concept of disease-related genes influencing response to treatment is not new. For example, variation in the gene coding for the 5-hydroxytryptamine 2A receptor has been associated with variation of clozapine response and increased susceptibility to schizophrenia [44,45]. Variation in the gene coding for the  $\beta$ -2-adrenergic receptor has been associated with airway responsiveness to  $\beta$ -2-receptor agonists and susceptibility to lower airway reactivity in patients with asthma [46,47]. Our results show that patients with a higher number of risk alleles were younger at the date of their first SU prescription. This may be the result of a more 'aggressive' form of T2DM. For many complex diseases such as T2DM, there may be multiple genetic backgrounds resulting in similar phenotypic disease, each requiring a different drug treatment. Our results support this concept, and support the use of disease-related genes in pharmacogenetic studies. We should emphasize, however, the fact that we have only begun to unravel the genetic determinants of drug response in T2DM and that although many of the genes are associated with  $\beta$ -cell function, the exact mechanism behind our finding remains unclear. Our results do provide some 'proof of principle' that the complex background of T2DM may ultimately result in the identification of different genetic subgroups of T2DM patients that require different pharmacotherapy. However, replication in an independent cohort and further elucidation of the causal mechanisms underlying SU response are warranted.

In conclusion, T2DM-associated risk alleles are associated with response to SU treatment in primary care T2DM patients. This suggests that individualization of T2DM treatment according to genetic profile may be an opportunity to improve clinical outcome.

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